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(54) 3-Aryl-4(3H)quinazolinone CCK antagonists and pharmaceutical formulations thereof.

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CHEMICAL ABSTRACTS, vol. 114, no. 19, May 13, 1991, Columbus, Ohio, USA YU, MELVIN J. et al. "Quinazolinone cholecystokinin-B receptor ligands" page 16, column 2, abstract-no. 277 878r

CHEMICAL ABSTRACTS, vol. 102, no. 5, February 4, 1985, Columbus, Ohio, USA AGARWAL, RAJESH et al. "Synthesis of 1-(2-arylin-dol-3-yl)-2-(3-(p-morpholino-phenyl)-6,8-disubstituted-quinazolin-4(3H)-on-2-yl)ethenes as CNS active and anti-inflammatory agents" page 570, column 1, abstract -no. 45 870z

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CHEMICAL ABSTRACTS, vol. 97, no. 1, July 5, 1982, Columbus, Ohio, USA AGARWAL, RAJESH et al. "Synthesis of 1-(2-aryl-1-indol-3-yl)-2-(3-(p-substituted-phenyl)-6-substituted-quinazolin-4-one-2-yl)ethenes as CNS active agents" page 606, column 2, abstract -no. 6 251g

CHEMICAL ABSTRACTS, vol. 105, no. 11, September 15, 1986, Columbus, Ohio, USAMAKOVEC, FRANCESCO et al. "New glutamic and aspartic derivatives with potent CCK-antagonistic activity" page 88, column 1, abstract- no. 91 509y

Description

This invention relates to biologically active quinazolinones. More particularly, this invention is directed to certain substituted quinazolinones which bind to receptors for cholecystokinin (CCK), e.g., those of the brain and pancreas, and to receptors for gastrin, e.g., those of the stomach. The compounds are CCK and gastrin antagonists and are useful in the treatment and prevention of CCK and gastrin-related disorders of the gastrointestinal, central nervous and appetite regulatory systems of warm-blooded vertebrates, especially humans.

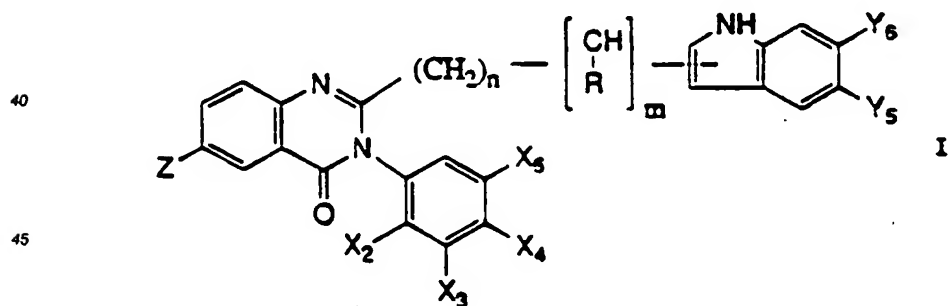
Cholecystokinin (CCK) is a neuropeptide found in both gastrointestinal tissue and the tissues of the central nervous system. CCK is believed to play an important role in appetite regulation. Among the effects of CCK are stimulation of colonic motility, stimulation of gall bladder contraction, stimulation of pancreatic enzyme secretion, and inhibition of gastric emptying. CCK reportedly coexists with dopamine in certain mid-brain neurons and thus may also play a role in the functioning of dopaminergic systems in the brain. Gastrin is a neuropeptide found particularly in the gastrointestinal tract. It is one of the primary natural stimulators of gastric acid secretion. It also has growth stimulatory effects on a variety of gastrointestinal tissues.

CCK and gastrin antagonists are useful in the treatment and prevention of CCK and gastrin-related disorders of the gastrointestinal and central nervous systems, as well as modulation of the appetite regulatory systems of warm-blooded vertebrates. The CCK/gastrin receptor family is thought to contain three receptor subtypes, for which the location of the prototype receptor is given in parentheses: CCK-A (pancreas), CCK-B (brain), and gastrin (stomach fundus).

Several classes of CCK receptor antagonists have been reported in the literature. One class comprises derivatives of cyclic nucleotides, for example, dibutyryl cyclic GMP. Another art recognized class of CCK antagonists comprise the C-terminal fragments and analogs of CCK. Another class of CCK receptor antagonists are amino acid derivatives including proglumide, a derivative of glutamic acid, and the N-acetyltryptophanes such as p-chlorobenzoyl-L-tryptophan. More recently certain substituted amino phenyl compounds were described as CCK antagonists in published European Patent Application 0166355. Because of the wide range of potential clinical applications of CCK binding compounds, intensive research efforts have been ongoing to define other compounds exhibiting CCK receptor binding properties.

This invention is directed to novel quinazolinones of Formula I below which have been found to exhibit CCK and gastrin antagonist activity. These compounds are useful in the treatment and prevention of CCK-related disorders of the gastrointestinal, central nervous, and appetite regulatory systems of mammals, especially humans. As gastrin antagonists, they are particularly useful in the treatment and prevention of gastrointestinal ulcers.

This invention is directed to compounds of the formula



in which

n is 1 or 2 and m is 0 or 1;

R is hydrogen, C₁-C₄ alkyl, benzyl, or phenyl;

Z is hydrogen or halo;

X₂, X₃, X₄, and X₅ are independently selected from the group consisting of hydrogen, halo, trifluoromethyl, C₁-C₆ alkoxy, C₁-C₆ alkyl, C₁-C₆ alkylthio, and -NR₂R₃, in which R₂ and R₃ are independently hydrogen, C₁-C₄ alkyl, benzyl, or phenyl, or R₂ and R₃ taken together with the nitrogen atom to which they are bonded form a 5- or 6-membered ring; or

X_r and X_{r+1}, in which r is 2, 3, or 4, taken together form a divalent C₃-C₅ alkylene group or

methylenedioxy; and

Y_5 and Y_6 are independently selected from the group consisting of hydrogen, C_1 - C_6 alkyl, C_1 - C_6 alkoxy, halo, and trifluoromethyl; and pharmaceutically acceptable salts thereof.

In the above formula, the term " C_1 - C_4 alkyl" means a straight or branched alkyl chain having from one to four carbon atoms. Such C_1 - C_4 alkyl groups are methyl, ethyl, *n*-propyl, isopropyl, *n*-butyl, isobutyl, *sec*-butyl, and *t*-butyl.

The term " C_1 - C_6 alkyl" means a straight or branched alkyl chain having from one to six carbon atoms or cycloalkyl having from three to six carbon atoms. Groups which are included in such term are methyl, ethyl, *n*-propyl, isopropyl, cyclopropyl, *n*-butyl, isobutyl, methylcyclopropyl, cyclobutyl, *sec*-butyl, *t*-butyl, *n*-pentyl, cyclopentyl, 2-methylbutyl, 3-methylbutyl, *n*-hexyl, cyclohexyl, 4-methylpentyl, and the like.

The term " C_1 - C_6 alkoxy" means a straight or branched chain alkyl group having from one to six carbon atoms or cycloalkyl having from three to six carbon atoms and joined through an oxygen atom. Groups which are included in such term are methoxy, ethoxy, *n*-propoxy, isopropoxy, cyclopropoxy, *n*-butoxy, isobutoxy, *t*-butoxy, *sec*-butoxy, cyclobutoxy, *n*-pentoxy, cyclopentoxy, 3-methylbutoxy, *n*-hexyloxy, 2-methylpentoxy, cyclohexyloxy, and the like.

The term " C_1 - C_6 alkylthio" means a straight or branched chain alkyl group having from one to six carbon atoms or cycloalkyl having from three to six carbon atoms and joined through a sulfur atom. Groups which are included in such term are methylthio, ethylthio, *n*-propylthio, isopropylthio, cyclopropylthio, *n*-butylthio, isobutylthio, *t*-butylthio, *sec*-butylthio, cyclobutylthio, *n*-pentylthio, cyclopentylthio, 2-methylbutylthio, *n*-hexylthio, 4-methylpentylthio, cyclohexylthio, and the like.

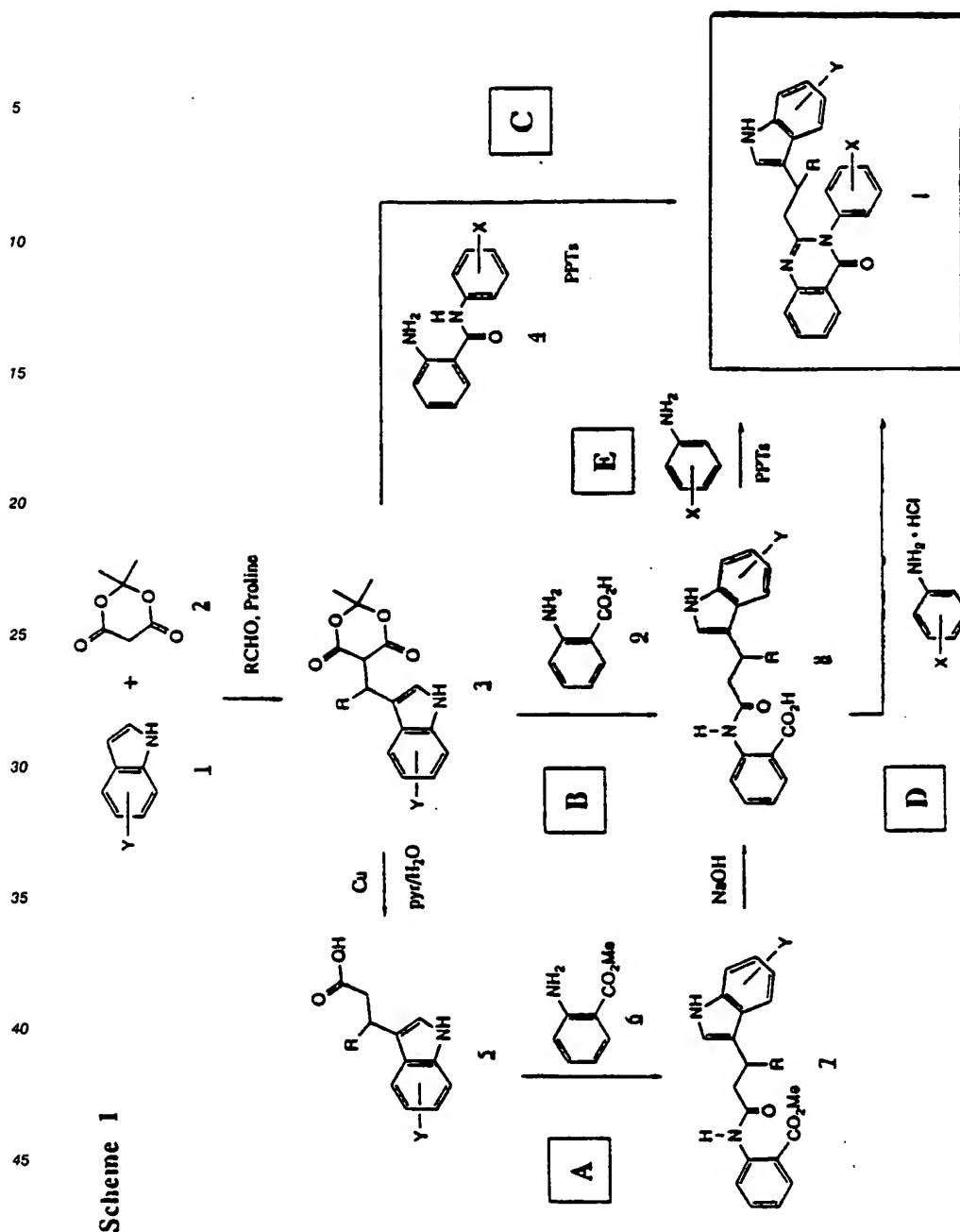
The term "halo" means any of fluoro, chloro, or bromo.

The term " C_3 - C_5 alkylene" means any of $-CH_2CH_2CH_2-$, $-CH_2CH_2CH_2CH_2-$, and $-CH_2CH_2CH_2CH_2CH_2-$.

One preferred group of compounds of Formula I are those in which m is 0, n is 2, and Z is hydrogen.

Another preferred group of the present compounds are those in which Y_6 is hydrogen and Y_5 is hydrogen, fluoro, chloro, or bromo. Of those preferred compounds, more preferred are those wherein X_2 and X_5 are hydrogen and at least one of X_3 and X_4 are hydrogen and the other is selected from the group consisting of C_1 - C_4 alkoxy, C_1 - C_6 alkyl or $-NR_2R_3$; most preferably, X_4 is hydrogen and X_3 is C_1 - C_4 alkoxy.

With reference to Scheme 1 following, the present compounds are generally prepared by reaction of indole 1 and Meldrum's acid 2 and an aldehyde RCHO (formaldehyde, $R=H$) in the presence of proline [Diane S. Farlow, Michael E. Flaugh, Sharon D. Horvath, Edward R. Lavagnino, and Paul Franc, *Organic Preparations and Procedures Int.* 13(1), 39 (1981)] to provide condensation adduct 3 which can be converted directly using Route C to quinazoline I by reaction with an *o*-amino-benzanilide 4 in the presence of pyridinium *p*-toluene-sulfonate (PPTS) in refluxing pyridine. Alternatively, via Route A condensation adduct 3 can be decarboxylated in aqueous pyridine in the presence of copper powder to provide the intermediate indolylpropionic acid 5. The latter reacts with a methyl anthranilate 6 to form ester 7. Hydrolysis of ester 7 provides the corresponding carboxylic acid 8 which reacts at elevated temperature with an aniline hydrochloride via Route D or with an aniline in the presence of PPTS [Route E] to provide quinazoline I. Intermediate acid 8 can be prepared directly from adduct 3 by reaction with an anthranilic acid 9 in refluxing pyridine [Route B].



50 The term "pharmaceutically acceptable salts" encompasses those salts that form by standard acid-base reactions with basic groups. Thus, the pharmaceutically acceptable salts of the present invention can be prepared by conventional chemical methods from those compounds of Formula I which contain a basic moiety. Generally, the salts are prepared by reacting the free base with a stoichiometric amount or with an excess of the desired salt-forming acid in a suitable solvent or combination of solvents. Suitable salt-forming acids include inorganic acids, such as hydrochloric, hydrobromic, sulfuric, sulfamic, phosphoric, nitric, and the like; organic acids, such as acetic, propionic, succinic, glycolic, stearic, lactic, citric, malic, tartaric, ascorbic, pantoic, maleic, hydroxymaleic, phenylacetic, glutamic, benzoic, salicylic, sulfanilic, 2-acetoxybenzoic, fumaric, toluenesulfonic, methane sulfonic, ethanesulfonic, oxalic, benzenesulfonic, picric, cinnamic,

and like acids.

The compounds of this invention bind to CCK receptors in the brain and/or peripheral sites such as the pancreas, gall bladder, and ileum. Their ability to antagonize CCK and gastrin makes these compounds useful as pharmaceutical agents for the treatment and prevention of disease states wherein CCK or gastrin may be involved, for example, gastrointestinal disorders, such as irritable bowel syndrome, ulcers, excess pancreatic or gastric secretion, acute pancreatitis, motility disorders, central nervous system disorders caused by CCK's interaction with dopamine, such as neuroleptic disorders, tardive dyskinesia, Parkinson's disease, psychosis or Gilles de la Tourette Syndrome, and disorders of appetite regulatory systems.

In another Embodiment of this invention there is provided pharmaceutical formulations comprising as an active ingredient an effective amount of a compound of Formula I and a pharmaceutically acceptable carrier, excipient, or diluent therefor. Such formulations can be prepared for oral or parenteral administration for the treatment and prevention of disorders of the gastrointestinal, central nervous, and appetite regulatory systems of warm-blooded vertebrates, especially man.

For oral use of an antagonist of CCK or gastrin of this invention, the selected compounds can be administered, for example, in the form of tablets or capsules, or as an aqueous solution or suspension. In the case of tablets, common excipients include binding agents, for example, syrup, acacia, gelatin, sorbitol, tragacanth, polyvinylpyrrolidone (Povidone), methylcellulose, ethylcellulose, sodium carboxymethylcellulose, hydroxypropylmethylcellulose, sucrose and starch; fillers and carriers, for example, corn starch, gelatin, lactose, sucrose, microcrystalline cellulose, kaolin, mannitol, dicalcium phosphate, sodium chloride and alginic acid; lubricants such as magnesium stearate; disintegrants such as croscarmellose, microcrystalline cellulose, corn starch, sodium starch glycolate and alginic acid; and suitable wetting agents such as lauryl sulfate. For oral administration in capsule form, useful diluents included lactose and dried corn starch. When aqueous suspensions are desirable for oral use, the active ingredient can be combined with emulsifying and suspending agents, for example, sorbitol, methylcellulose, glucose/sugar syrup, gelatin, hydroxyethylcellulose, carboxymethylcellulose, aluminum stearate gel, or hydrogenated edible oils, for example, almond oil, fractionated coconut oil, oily esters, propylene glycol, or ethyl alcohol; flavoring agents such as peppermint, oil of wintergreen, cherry flavoring, and the like; and preservatives, such as methyl or propyl p-hydroxybenzoates and ascorbic acid.

The pharmaceutical formulations in accordance with this invention can also be prepared for parenteral use. Such formulations typically take the form of sterile isotonic solutions of the active ingredient according to standard pharmaceutical practice.

The appropriate dose of the compound of the present invention for its use as an antagonist of CCK or gastrin in humans will vary according to the age, weight and response of the individual patient, as well as the severity of the patient symptoms and the nature of the condition being treated. Thus, the preferred daily dose will normally be determined by the prescribing physician. However, in most instances, effective daily doses of the compounds of this invention will range from about 0.05 mg/kg to about 50 mg/kg and preferably from about 0.5 mg/kg to about 20 mg/kg in a single or in divided doses.

The following examples are provided to further illustrate the compounds of this invention and the methods for their preparation.

Example 1

2-[2-(3-Indolyl)ethyl]-3-phenyl-4-(3H)quinazolinone [Route A/E].

To a solution of 3-(3-indolyl)propionic acid (6.0 g, 32 mmol) in 100 ml THF at room temperature was added 1,1-carbonyldiimidazole (5.14 g, 32 mmol). The reaction mixture was stirred under a dry atmosphere for 30 minutes, after which methyl anthranilate (4.79 g, 32 mmol) was added, and the reaction mixture was stirred at reflux for 30 minutes. No reaction was detected by thin layer chromatography (TLC). Pyridinium p-toluene-sulfonate (PPTS) [6.36 g, 25 mmol] was added, and the mixture was stirred at reflux for two days. The reaction mixture was allowed to cool, and the solvent was removed in vacuo. The resulting product was taken up in ethyl acetate and washed (1N HCl, H₂O, satiated NaHCO₃, and brine) and dried over anhydrous magnesium sulfate. Concentration in vacuo provided a light yellow solid which, when triturated with ethyl acetate/hexane, gave, upon filtration, 7.19 g (70% yield) of 3-(3-indolyl)-N-(2-methoxycarbonylphenyl)-propionamide as an off white granular solid.

The methyl ester product (7.19 g, 22.3 mmol) was dissolved in 75 ml methanol and 25 ml 1N sodium hydroxide. The reaction mixture was stirred at reflux for 30 minutes, cooled to room temperature, and concentrated in vacuo. The aqueous residue was diluted with water and washed once with diethyl ether. The aqueous phase was separated, acidified with 40 ml of 1N HCl, and extracted with ethyl acetate. The

organic layer was washed with brine and dried over anhydrous sodium sulfate. Evaporation to dryness in vacuo provided a solid which, upon trituration with 40% ethyl acetate/hexanes and filtration, provided 6.34 g (92% yield) of an off-white, granular solid, the physical chemical data for which [MS 308 (M⁺)] was consistent with that expected for the intermediate 3-(3-indolyl)-N-(2-carboxyphenyl)propionamide.

5 To a solution of 3-(3-indolyl)-N-(2-carboxyphenyl)propionamid (6.20 g, 20 mmol) in 50 ml of THF at room temperature was added 1,1-carbonyldiimidazole (3.26 g, 20 mmol). The reaction mixture was stirred under a dry atmosphere for 30 minutes after which aniline (2 ml, 22 mmol) and PPTS (4.03 g, 16 mmol) were added. The resulting reaction mixture was stirred under reflux for two days. The reaction mixture was allowed to cool and evaporated to dryness in vacuo. The product residue was taken up in ethyl acetate and
10 washed (1N HCl, water, saturated NaHCO₃, and brine), dried over anhydrous sodium sulfate, and concentrated in vacuo to a brown oil which crystallized from ethyl acetate/hexanes. The product was recrystallized from methanol/ethyl acetate to provide 2.28 g of 2-[2-(3-indolyl)ethyl]-3-phenyl-4-(3H)-quinazolinone as white fluffy needles; m.p. 193-194 °C; FABMS and EIMS 366 (M⁺); IR (KBr) 1672 CM⁻¹.

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Analysis, calculated for C ₂₄ H ₁₉ N ₃ O:			
Calc.:	C, 78.88;	H, 5.24;	N, 11.50;
Found:	C, 78.65;	H, 5.26;	N, 11.36.

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Example 2

2-(3-Indolylmethyl)-3-phenyl-4-(3H)quinazolinone.

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[Route A/E modified] To a solution of 3-indoleacetic acid (5.79 g, 33.1 mmol) in 100 ml THF at room temperature was added 1,1-carbonyldiimidazole (5.36 g, 33.1 mmol). The reaction mixture was stirred for 15 minutes under nitrogen. Methyl anthranilate (5.0 g, 33.1 mmol) and PPTS (20.0 g, 80 mmol) were added to the reaction mixture which was then stirred at reflux for 24 hours, cooled, and concentrated to near dryness in vacuo. The product residue was partitioned between 1N hydrochloric acid and ethyl acetate. The two-
30 phase mixture was heated until two clear layers were obtained. The aqueous layer was separated and the organic layer was allowed to cool, washed (water, 1N NaOH, and brine), dried over anhydrous sodium sulfate, and concentrated in vacuo to dryness. The resulting product was triturated with ethyl acetate/hexane and collected by vacuum-assisted filtration to give 8.85 g of 2-(3-indolyl)-N-(2-methoxycarbonylphenyl)-acetamide as a pale yellow granular solid which was deesterified by refluxing it in a mixture of 100 ml of
35 methanol and 32 ml of 1M NaOH for 30 minutes. The product acid was isolated by concentrating the deesterification mixture in vacuo, acidifying with 40 ml of 1N hydrochloric acid, and then extracting it into ethyl acetate. The organic layer was washed with brine, dried over anhydrous sodium sulfate, and concentrated in vacuo to dryness. Trituration with ethyl acetate/hexanes and vacuum-assisted filtration provided 8.0 g of 2-(3-indolyl)-N-(2-carboxyphenyl)acetamide as an off-white solid: mp 205-207 °C.

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Analysis, calculated for C ₁₇ H ₁₄ N ₂ O ₃ :			
Calc.:	C, 69.38;	H, 4.79;	N, 9.52;
Found:	C, 69.29;	H, 4.93;	N, 9.45.

45

To a solution of the above product (7.0 g, 23.8 mmol) in 100 ml THF at room temperature was added 1,1-carbonyldiimidazole (4.24 g, 26.2 mmol). The reaction mixture was stirred for 30 minutes after which aniline (2.4 ml, 26 mmol) and PPTS (15.8 g, 63 mmol) were added. The reaction mixture was stirred at
50 reflux for 2 days under nitrogen, cooled, and concentrated to near dryness in vacuo. The residue was partitioned between ethyl acetate and 1N hydrochloric acid. The organic phase (which contained an undissolved solid) was washed once with 1N hydrochloric acid and 3 times with water. The organic phase was separated and evaporated in vacuo. The residue was taken up in toluene and again concentrated to near dryness. The resulting solid was triturated with ethyl acetate/hexanes and collected by vacuum-
55 assisted filtration. The product was then dispersed in methanol/ethyl acetate, heated to boiling, and allowed to cool. Vacuum-assisted filtration provided 3.08 g of 2-(3-indolyl)-N-(2-anilinoacetylphenyl)acetamide as whit needles: mp 235-236.5 °C; FDMS: 369 (M⁺).

Analysis, calculated for C ₂₃ H ₁₉ N ₃ O ₂ :			
Calc.:	C, 74.78;	H, 5.18;	N, 11.37;
Found:	C, 75.15;	H, 5.10;	N, 11.33.

A mixture of 500 mg of the above product, and 26 mg of p-toluenesulfonic acid in 20 ml of toluene was refluxed under drying tube with azeotropic removal of water for 3 hours. Additional p-toluenesulfonic acid (26 mg) was added to the reaction mixture before it was refluxed for an additional 21 hours. The mixture was cooled, diluted with ethyl acetate, washed (saturated NaCO₃ and brine), dried over anhydrous sodium sulfate, and concentrated in vacuo to near dryness. The residue was triturated with methanol/ethyl acetate and filtered. The filtrate was concentrated in vacuo and chromatographed (50% ethyl acetate/hexanes, SiO₂) to give the titled product as a yellow oil which crystallized from ethyl acetate/hexanes as fine white needles (150 mg): mp 191-192 °C; EIMS: 351 (M⁺).

Analysis, calculated for C ₂₃ H ₁₇ N ₃ O:			
Calc.:	C, 78.61;	H, 4.88;	N, 11.96;
Found:	C, 78.79;	H, 4.96;	N, 12.02.

Example 3

2-[2-(5-Methoxyindol-2-yl)ethyl]-3-phenyl-4-(3H)quinazolinone [Route A/D].

A solution of 5-methoxyindole (10.0 g, 67.9 mmol), 5.5 ml formaldehyde (37% aqueous solution), Meldrum's acid (10 g, 69 mmol), and proline (0.4 g) in 40 ml of acetonitrile was stirred at room temperature for 16 hours. The reaction mixture was concentrated in vacuo to provide a brown foam. The product was taken up in about 50 ml of acetone, and about 50 ml of water was added to the cloud point. The mixture was allowed to stand in the freezer and, after crystallization began, additional water was added with swirling again to the cloud point. Vacuum-assisted filtration provided 16.2 g of 2-(2,2-dimethyl-4,6-dioxo-1,3-dioxan-5-yl)methyl-5-methoxyindole as sand-colored crystals.

The product (16.2 g, 53.4 mmol), copper powder (390 mg), and water (15 ml) in 140 ml of pyridine was refluxed for 3 hours. The reaction mixture was cooled, filtered, and concentrated in vacuo to near dryness. The product was taken up in toluene and the resulting mixture again concentrated in vacuo to dryness. The residue was then dissolved in one liter of diethyl ether, washed (500 ml of 1N hydrochloric acid, 500 ml 20% aqueous ammonium chloride, water, and brine), dried over anhydrous sodium sulfate, and concentrated in vacuo to dryness. Crystallization of the product from chloroform/hexane yielded 9.94 g (85%) of 3-(5-methoxyindol-3-yl)propionic acid as a brown powder: mp 123-128 °C.

To a solution of of the foregoing product (6.95 g, 31.7 mmol) in 100 ml THF at room temperature under a drying tube was added 1,1-carbonyldiimidazole (5.14 g, 31.7 mmol). The reaction mixture was stirred for one hour after which methyl anthranilate (4.79 g, 31.7 mmol) and PPTS (7.96 g, 31.7 mmol) were added. The reaction mixture was stirred at reflux for three days after which an additional 7.96 g of PPTS was added, and reflux was continued for 2 days. The reaction mixture was cooled, diluted with ethyl acetate, washed (1N hydrochloric acid, water, saturated sodium bicarbonate, and brine), dried over anhydrous sodium sulfate, and concentrated in vacuo to dryness. Crystallization of the resulting residue from ethyl acetate/hexanes provided 5.21 g (47%) of 3-(5-methoxyindol-3-yl)-N-(2-methoxycarbonylphenyl)-propionamide.

The foregoing propionamide ester (5.2 g) was dissolved in a mixture of 50 ml of methanol and 16.5 ml of 1N sodium hydroxide solution, and the mixture was refluxed for 30 minutes, cooled, and concentrated in vacuo to near dryness. The product residue was diluted with water and washed once with ether. The aqueous layer was acidified with 20 ml of 1N hydrochloric acid and extracted with ethyl acetate. The organic layer was separated, washed with brine, dried over anhydrous sodium sulfate, and concentrated in vacuo to dryness. Crystallization of the product from ethyl acetate/hexane provided 4.53 g (91% yield) of 3-(5-methoxyindol-3-yl)-N-(2-carboxyphenyl)propionamide as a brown powder: mp 168-170 °C; FDMS: 338 (M⁺).

Analysis, calculated for C ₁₉ H ₁₈ N ₂ O ₄ :			
Calc.:	C, 67.45;	H, 5.36;	N, 8.28;
Found:	C, 67.26;	H, 5.63;	N, 8.07.

5

To a solution of 3-(5-methoxyindol-3-yl)-N-(2-carboxyphenyl)propionamide (4.0 g, 11.8 mmol) in 50 ml of THF at room temperature was added 1,1-carbonyldiimidazole (1.92 g, 11.8 mmol). The reaction mixture was stirred for one hour after which aniline hydrochloride (3.06 g, 23.6 mmol) and 25 ml of dimethyl-formamide were added. The reaction mixture was stirred at reflux under a drying tube for two days, allowed to cool, and concentrated in *vacuo* to dryness. The residue was partitioned between 1N hydrochloric acid and ethyl acetate. The organic layer was washed (water, 1N sodium hydroxide, and brine), dried over anhydrous sodium sulfate, and concentrated in *vacuo* to dryness. Crystallization of the product from ethyl acetate/trace methanol provided 1.86 g of 2-[2-(5-methoxyindol-3-yl)-ethyl]-3-phenyl-4-(3H)quinazolinone as a white powder: mp 220-221 °C; EIMS: 395 (M⁺).

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Analysis, calculated for C ₂₅ H ₂₁ N ₃ O ₂ :			
Calc.:	C, 75.93;	H, 5.35;	N, 10.63;
Found:	C, 75.65;	H, 5.30;	N, 10.35.

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The foregoing intermediate product 3-(5-methoxyindol-3-yl)-N-(2-carboxyphenyl)propionamide can be prepared directly from the corresponding Meldrum's Adduct [3-(2,2-dimethyl-4,6-dioxo-1,3-dioxane-5-yl)-methyl-5-methoxyindole] (4.41 g, 14.5 mmol) by reaction with an equivalent amount of anthranilic acid (1.99 g, 14.5 mmol) in 30 ml of pyridine. The reaction is carried out at reflux for two hours after which the pyridine is azeotropically removed under vacuum with toluene [Route B].

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Example 4

2-[2-(5-Bromoindol-3-yl)ethyl]-3-(3-isopropoxyphenyl)-4-quinazolinone [Route C].

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3-Nitrophenol (50.0 g, 360 mmol), isopropyl iodide (76.19 g, 450 mmol), and potassium carbonate (60 g) were combined and refluxed under nitrogen overnight in 400 ml of acetone. The reaction mixture was cooled, and the solvent was removed in *vacuo*. The reaction was combined with about 300 ml of water and thereafter extracted with four 100 ml portions of ethyl acetate. The ethyl acetate extracts were combined and washed twice with 1N sodium hydroxide and brine, dried over anhydrous sodium, and evaporated to provide 56 g (86%) of 3-isopropoxynitrobenzene as a clear yellow oil.

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A solution of 3-isopropoxynitrobenzene (8.5 g, 50 mmol) and PtO₂ (0.3 g) in 200 ml of ethanol was shaken under 40 psi hydrogen at room temperature for 1.5 hours. The mixture was filtered through Celite, and the filtrate was concentrated in *vacuo* to provide 7.08 g of light oil (3-isopropoxyaniline). The oil was added to isatoic anhydride (7.35 g, 45 mmol) along with 15 ml of ethyl acetate. The mixture was heated in a 90 ° oil bath under nitrogen for two hours. The product crystallized from the reaction mixture upon the addition of hexanes. Filtration of the reaction mixture provided 10.19 g (83%) of 2-amino-N-(3-isopropoxyphenyl)benzamide as a white solid.

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A solution of 5-bromoindole (10.07 g, 51 mmol), Meldrum's acid (7.38 g, 51 mmol), proline (1.24 g), and 5.2 ml of 30% aqueous formaldehyde in 100 ml of acetonitrile was allowed to stand at room temperature for 24 hours. The acetonitrile was removed in *vacuo*. Crystallization of the product from methanol (50 ml) provided 15.83 g of 3-(2,2-dimethyl-4,6-dioxo-1,3-dioxane-5-yl)methyl-5-bromoindole as a white solid.

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A solution of 3-(2,2-dimethyl-4,6-dioxo-1,3-dioxane-5-yl)methyl-5-bromoindole (4.12 g, 12 mmol), 2-amino-N-(3-isopropoxyphenyl)benzamide (3.48 g, 13 mmol), and PPTS (1.64 g, 6.5 mmol) in 50 ml of pyridine was refluxed for 3.5 days. The reaction mixture was then evaporated in *vacuo* to dryness, and the residue was taken up in methylene chloride. The product was chromatographed (30% ethyl acetate/hexanes, SiO₂) and the title product (2.13 g, 36%) crystallized by allowing the fractions containing product to evaporate: mp 179-181 °C.

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55

Analysis, calculated for $C_{27}H_{24}N_3O_2Br$:			
Calc.:	C, 64.55;	H, 4.81;	N, 8.36;
Found:	C, 64.79;	8, 5.01;	N, 8.36.

5

Example 5

10 2-[2-Phenyl-2-(3-indolyl)ethyl]-3-(3-methoxyphenyl)-4-(3H)quinazolinone [Route C modified].

Indole (10 g, 85 mmol), Meldrum's acid (12.3 g, 85 mmol), benzaldehyde (18.11 g, 171 mmol) and proline (0.05 g) were combined with 50 ml of acetonitrile and stirred in an oil bath at approximately 35-40 °C for two hours [Y. Oikawa, H. Hirasawa, and O. Honemitsu, Tetrahedron Letters, 1759 (1978)]. The reaction mixture was concentrated in vacuo to dryness. The residue was slurried with methanol and filtered to provide 19.99 grams of 3-[α -(2,2-dimethyl-4,6-dioxo-1,3-dioxane-5-yl)benzyl]indole. A portion of the product was recrystallized from ethyl acetate to provide a white crystalline product: mp 147-150 °C.

20

Analysis, calculated for $C_{21}H_{19}NO_4$:			
Calc.:	C, 72.19;	H, 5.48;	N, 4.01;
Found:	C, 72.19;	H, 5.61;	N, 4.08.

Isatoic anhydride (32.63 g, 200 mmol) and 3-methoxyaniline (24.63 g, 200 mmol) were combined neat and heated at 120 °C for 2 hours. The reaction product was taken up in methylene chloride and chromatographed (20% ethyl acetate/hexanes, SiO_2), to provide 38.75 g (80%) of 2-amino-N-(3-methoxyphenyl)benzamide. An analytical sample was obtained by recrystallization from ethyl acetate: mp 75-77 °C.

30

Analysis, calculated for $C_{14}H_{14}N_2O_2$:			
Calc.:	C, 69.40;	H, 5.82;	N, 11.56;
Found:	C, 69.63;	H, 5.84;	N, 11.60.

2-Amino-N-(3-methoxyphenyl)benzamide (2.77 g, 11.5 mmol), 3-[α -(2,2-dimethyl-4,6-dioxo-1,3-dioxane-5-yl)benzyl]indole (4.00 g, 11.5 mmol), and PPTS (1.44 g, 5.7 mmol) in 20 ml of pyridine was refluxed for seven days. The reaction mixture was concentrated in vacuo to an oil and partitioned between ethyl acetate and water. The ethyl acetate layer was separated, washed (1N hydrochloric acid, saturated sodium bicarbonate, and brine), dried over anhydrous sodium sulfate, and concentrated in vacuo to dryness. The reaction was incomplete. The product was therefore mixed with PPTS (3 g) and 10 ml of 2,4,6-collidine and refluxed for five hours. The product was worked up as above to provide an oil which, upon standing for about one month, began to crystallize. Trituration with ethyl acetate and filtration provided 1.4 g of the title product: mp 160-164 °C.

45

Analysis, calculated for $C_{31}H_{25}N_3O_2 \cdot 1/3 C_4H_8O_2$:			
Calc.:	C, 77.53;	H, 5.57;	N, 8.39;
Found:	C, 77.58;	H, 5.45;	N, 8.47.

50

Test Procedures for CCK And Gastrin Receptor Binding (IC_{50})Brain

55 Brain CCK receptor binding was performed using mouse brain membranes according to the method of Chang and Lotti (Proc. Natl. Acad. Sci. 83: 4923-4926, 1986). Male CF-1 mice, 23-25 g were sacrificed by cervical dislocation, the forebrain removed and placed in ice cold 50 mM Tris buffer, pH 7.4. The tissue was homogenized in 100 volumes of the Tris buffer with a Brinkman Polytron or T kmar Tisumizer and then

centrifuged at 40,000 g for 10 min. Pellets were resuspended in Tris buffer, centrifuged as above and then resuspended in 100 volumes of assay buffer, pH 6.5 (20 mM N-2-hydroxyethyl-piperazine-N'-2-ethane sulfonic acid (HEPES), 1 mM ethylene glycol bis(2-aminoethyl ether-N,N,N',N'-tetraacetic acid) (EGTA), 5 mM MgCl₂, 130 mM NaCl, and 0.25 mg/ml bacitracin). The binding assay consisted of 50 μ L of compound (or buffer for total binding), 50 μ L of ¹²⁵I-CCK-8 sulfate (20 pM) (Amersham IM-159), 200 μ L of assay buffer and 200 μ L of homogenate (80-120 μ g protein). The samples were incubated at room temperature (25°) for 2 hours, and they were then filtered through GF/B glass fiber filters (soaked in wash buffer for 2 hours before use) using a 48 well Brandel cell harvester designed for receptor binding. The filters were washed twice with 3 ml of 50 mM Tris buffer, pH 7.4, containing 0.01% BSA and then counted for radioactivity in plastic tubes with a Micromedic 10/600 automatic gamma counter.

Compounds were dissolved in dimethyl sulfoxide (DMSO) at a concentration of 10 mM and then further diluted with assay buffer. The concentration of DMSO in the incubation was 0.1% or less and had no effect on the assay at that level. IC-50 values of displacement curves were determined using 7 concentrations of compound and were calculated using the ALLFIT computer program of DeLean, Munson and Rodbard (*Am. J. Physiol.* 235: E97-E102, 1978). Non-specific binding was determined as the displacement of the radioligand by 100 nM CCK-8 sulfate.

Pancreas

Binding to peripheral type CCK receptors in rat pancreas was done according to the method of Chang et al. (*Mol. Pharmacol.* 30: 212-217, 1986) using ³H-L364, 718. Pancreas was obtained from male Sprague-Dawley rats, 150-200 g, after decapitation, and dissected free from adipose and connective tissue. The tissue was homogenized in 30 volumes of 50 mM Tris buffer, pH 7.4 and centrifuged at 40,000 g for 10 min. The tissue pellet was washed by resuspension and centrifugation as described above. The final pellet was suspended in 500 volumes of assay buffer (50 mM Tris buffer, pH 7.4, 5 mM MgCl₂, 0.14 mg/ml bacitracin, and 5 mM dithiothreitol) to give a protein concentration of 30-60 μ g/200 μ L. Reagent volumes for the assay were the same as those used for CCK binding to brain membranes. Tritium labeled L-364,718 (Dupont NEN, NET-971) was used as the ligand at a concentration of 0.4-0.6 nM. The samples were incubated 1 hour at room temperature and then filtered as described for the CCK-brain receptor. Scintillation cocktail was added to the filters which were counted for radioactivity using a Micromedic Taurus automatic liquid scintillation counter.

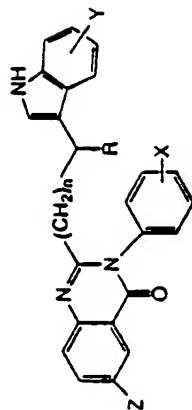
Compound samples were prepared and IC-50 values were determined as described for the CCK-brain experiments. Non-specific binding was that amount left bound to the filters after adding 100 nM L-364,718.

Gastric Mucosa

The method used for gastrin binding to guinea pig stomach mucosal membranes was similar to that described by Takeuchi, Speir and Johnson (*Am. J. Physiol.* 237(3): E284-E294, 1979). Guinea pig stomach fungus was obtained from male Hartley guinea pigs, 300-350 g, and the mucosa was scraped off with a glass slide. The mucosa was homogenized in 50 mM Tris buffer, pH 7.4, containing 1 mM phenylmethanesulfonyl fluoride using a Dounce glass homogenizer, and the suspension was centrifuged at 40,000 g for 10 min. The resulting pellet was resuspended and centrifuged once more, the final pellet was then suspended in 100 ml assay buffer per 1 guinea pig stomach to give a protein concentration of 200-300 μ g/200 μ L. The assay buffer consisted of 50 mM Tris buffer, pH 7.4, 5 mM MgCl₂, 0.14 mg/ml bacitracin, and 1 μ g/ml each of leupeptin, chymostatin, aprotinin and pepstatin. Reagent volumes for the assay were the same as those used for CCK binding to brain membranes. The radioactive ligand was 20 pM ¹²⁵I-gastrin I, from DuPont NEN (NEX-176). The samples were incubated 3 hours at room temperature and filtered and counted as described for CCK binding to brain membranes. Compound samples were prepared and IC-50 values were determined as described for the CCK-brain receptor binding. Non-specific binding was determined using 100 nM gastrin I (human synthetic from Sigma Chemical Co.).

Physical data, receptor binding data, and preparative methods are provided in Table I following for the foregoing Examples 1-5 as well as additional Examples 6-66. The compound of each example is identified by reference to the structural formula preceding each group of examples. The method for preparing each compound is indicated by reference to Procedure A-E corresponding to the procedures identified in the foregoing Examples 1-5 and Scheme 1.

Table I
Melting Point, Preparative Method, and CCK and Gastrin Receptor Binding Data



Example	X	Y	Z	n	R	m.p. (°C)	Proc- edure	Brain ¹	Pancreas ¹	Gastrin ¹
1	H	H	H	1	H	193-194	A, E	0.67±0.15 ⁷	378 (10μM)	
2	H	H	H	0	H	191-192	A, E ⁸	368 (10μM)		
3	H	5-MeO	H	1	H	220-221	A, D	0.41		
4	3-IPrO	5-Br	H	1	H	179-181	C	0.0093±0.0015 ⁷	-1.48(10μM)	0.16±0.04 ⁷
5	3-MeO	H	H	1	Ph	160-164	C ⁸	578(1μM)		

Table I (Cont.)
Melting Point, Preparative Method, and CCK and Gastrin Receptor Binding Data

Example	X	Y	Z	n	R	m.p. (°C)	Proc- edure	Brain ¹	Pancreas ¹	Gastrin ¹
6	H	5-Cl	H	1	H	228-229.5	B, D	85%, 23% (1µM)		
7	H	6-Cl	H	1	H	114-117	A, D	1.1		
						(dec)				
8	2-F	H	H	1	H	137-139	A, E ²	75% (10µM)		
9	2-Cl	H	H	1	H	182-185	A, E ²	44% (10µM)		
10	2-MeO	H	H	1	H	161-162	A, E ²	74% (10µM)		
11	2-CF ₃	H	H	1	H	149-150	A, E ²	35% (10µM)		
12	3-F	H	H	1	H	165.5-167.5	C	0.73±0.17 ⁷	11% (10µM)	
13	3-Cl	H	H	1	H	164-166	A, E ³	0.69±0.16 ⁷	17% (10µM)	
14	3-Me	H	H	1	H	175-177	C	0.15±0.01 ⁷	0.6% (10µM)	
15	3-Br	H	H	1	H	163-165	C	0.37±0.03 ⁷	-5.7% (10µM)	
16	3-Br	5-Cl	H	1	H	209-211	C	0.19	2% (10µM)	

Table I (Cont.)
Melting Point, Preparative Method, and CCK and Gastrin Receptor Binding Data

Example	X	Y	Z	n	R	m.p. (°C)	Proc- edure	Brain ¹	Pancreas ¹	Gastrin ¹
17	3-MeO	H	H	1	H	151-152	A, E ²	0.16±0.03 ⁷	168(10μM)	
18	3-MeO	5-F	H	1	H	229-231	C	0.11±0.01 ⁷		
19	3-MeO	5-Cl	H	1	H	234-236	C	0.047±0.003 ⁷		
20	3-MeO	5-Br	H	1	H	219-221	C	0.038±0.003 ⁷	5.38(10 μM)	1.3±0.4 ⁷
21	3-MeO	5-Me	H	1	H	169-171	C	0.055±0.003 ⁷		
22	3-MeO	5-MeO	H	1	H	166-168	C	0.067±0.005 ⁷		
23	3-EtO	5-Br	H	1	H	224-226	C	0.034±0.007 ⁷	-148(10μM)	
24	3-EtO	5-MeO	H	1	H	204-206	C	0.033		
25	3-IPrO	H	H	1	H	158-161	C	0.026±0.0003 ⁷		
26	3-IPrO	5-Cl	H	1	H	179-181	C	0.019±0.005 ⁷		
27	3-IPrO	5-MeO	H	1	H	166-168	C	0.019		

Table I (Cont.)
Melting Point, Preparative Method, and CCK and Gastrin Receptor Binding Data

Example	X	Y	Z	n	R	m.p. (°C)	Proc- edure	Brain ¹	Pancreas ¹	Gastrin ¹
28	3-Et	H	H	1	H	157-159	B, D ⁴	0.072±0.001 ⁷	2.63(10 μ M)	
29	3-Et	5-Br	H	1	H	166-168	C	0.046±0.010 ⁷	-4.58(10 μ M)	1.1±0.2 ⁷
30	3-Et	5-Cl	H	1	H	157-160	C	0.030	-9.18(10 μ M)	
31	3-Et	5-MeO	H	1	H	195-197	C	0.022	118(10 μ M)	
32	3-MeS	5-Br	H	1	H	183-185	C	0.046±0.008 ⁷		
33	3-MeS	5-MeO	H	1	H	138-140	C	0.034		
34	3-CF ₃	5-Br	H	1	H	175-177	C	0.23±0.03 ⁷		
35	3-CF ₃	5-Cl	H	1	H	154-158	C	0.23		
36	3-CF ₃	5-Me	H	1	H	152-155	C	0.19		
37	3-CF ₃	5-MeO	H	1	H	218-220	C	0.13		

Table I (Cont.)
Melting Point, Preparative Method, and CCK and Gastrin Receptor Binding Data

Example	X	Y	Z	n	R	m.p. (°C)	Proc- edure	Brain ¹	Pancreas ¹	Gastrin ¹
38	3-NMe ₂	5-Br	H	1	H	242-244	C	0.016±0.001 ⁷		
39	3-NMe ₂	5-Cl	H	1	H	225-227	C	0.013		
40	3-pyrrolidino	5-Br	H	1	H	250-251	C	0.022±0.003 ⁷		
41	3,5-diMeO	H	H	1	H	224-225	A, 1 ²	1.8		
42	3,4-OCH ₂ OH	H	H	1	H	189-191	A, 1 ²	0.19±0.04 ⁷		
43	4-MeO, 3-nPr	5-Br	H	1	H	221-223	C	85% (1μM)		
44	4-EtO, 3-Me	5-Br	H	1	H	215-217	C	0.065		
45	3-Me, 4-MeO	5-Br	H	1	H	241-243	C	0.029		
46	3,4-diMeO	5-Br	H	1	H	226-229	C	0.13±0.03 ⁷		
47	4-Et	5-Br	H	1	H	250-251	C	0.028±0.004 ⁷		
48	4-iPr	5-Br	H	1	H	238-240	C	0.037±0.013 ⁷		
49	4-iPr	5-Cl	H	1	H	240-241	C	0.038		
50	4-nBu	5-Br	H	1	H	238-240	C	0.93		

Table I (Cont.)
Melting Point, Preparative Method, and CCK and Gastrin Receptor Binding Data

Example	X	Y	Z	n	R	m.p. (°C)	Proc- edure	Brain ¹	Pancreas ¹	Gastrin ¹
51	4-MeO	H	H	1	H	229-230	A, E ²	0.098±0.007 ⁷		
52	4-MeO	5-Br	H	1	H	252-253	C	0.031±0.006 ⁷	0.3% (10µM)	
53	4-EtO	5-Br	H	1	H	253-256 (dec)	C	0.088±0.010 ⁷		
54	4-nPrO	5-Br	H	1	H	253-255 (dec)	C	1.2		
55	4-iPrO	5-Br	H	1	H	240-242 (dec)	C	0.11±0.02 ⁷		
56	4-nBuO	5-Br	H	1	H	256-258	C	1.1		
57	4-NMe ₂	5-Br	H	1	H	285-287 (dec)	C	0.033±0.006 ⁷		
58	4-NMe ₂	5-Cl	H	1	H	283-286 (dec)	C	0.087		
59	4-NEt ₂	5-Br	H	1	H	247-249 (dec)	C	0.047		

Table I (Cont.)
Melting Point, Preparative Method, and CCK and Gastrin Receptor Binding Data

Example	X	Y	Z	n	R	m.p. (°C)	Proc- edure	Brain ¹	Pancreas ¹	Gastrin ¹
60	4-MeS	5-Br	H	1	H	257-259	C	0.037±0.010 ⁷		
61	4-CF ₃	H	H	1	H	227-229	A, D ³	0.538(10μM)		
62	H	H	H	2	H	202-204	A, E	208(10μM)	118(10μM)	
63	3-MeO	5-Br	H	1	Me	208-211	C	0.10		1.6±0.1 ⁷
64	3-Et	5-Br	H	1	Me	186-188	C	0.10±0.01 ⁷		
65	3-MeO	H	H	1	φCH ₂	foam	C	708(10μM)		
66	3-IPrO	5-Br	Cl	1	H	180-185	C	908(1μM)		

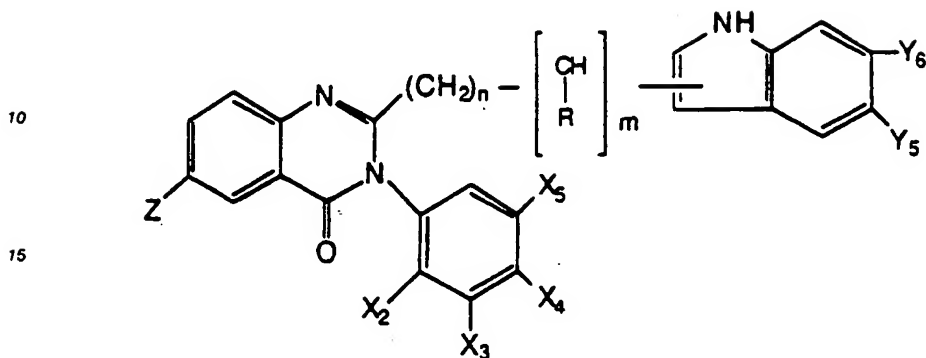
1. IC₅₀ (μM, Mean±SEM) or % Inhibition at specified concentration in binding assay. Values without SEM were obtained for n=1.
2. Standard procedure except used DME as solvent in final reaction.
3. Standard procedure except used DME/DMF(50:30) as solvent in final reaction.
4. Standard procedure except used DME/DMF (1:1) as solvent in final reaction.
5. Final closure to the quinazolinone ring required refluxing the amide intermediate with TsOH in toluene with azeotropic removal of water.
6. Standard procedure except used collidine as solvent in final reaction.
7. n=3

Claims

Claims for the following Contracting States : DE, FR, GB, IT

1. A compound of the formula

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in which

n is 1 or 2 and m is 0 or 1;

R is hydrogen, C₁-C₄ alkyl, benzyl, or phenyl;

Z is hydrogen or halo;

25

X₂, X₃, X₄, and X₅ are independently selected from the group consisting of hydrogen, halo, trifluoromethyl, C₁-C₆ alkoxy, C₁-C₆ alkyl, C₁-C₆ alkylthio, and -NR₂R₃, in which R₂ and R₃ are independently hydrogen, C₁-C₄ alkyl, benzyl, or phenyl, or R₂ and R₃ taken together with the nitrogen atom to which they are bonded form a 5- or 6-membered ring; or

30

X_r and X_{r+1}, in which r is 2, 3, or 4, taken together form a divalent C₃-C₅ alkylene group or methylenedioxy; and

Y₅ and Y₆ are independently selected from the group consisting of hydrogen, C₁-C₆ alkyl, C₁-C₆ alkoxy, halo, and trifluoromethyl; and pharmaceutically acceptable salts thereof.

2. A compound of Claim 1, in which Z is hydrogen.

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3. A compound of Claim 2, in which m is 0 and n is 2.

4. A compound of Claim 3, in which Y₅ is hydrogen, C₁-C₆ alkyl, C₁-C₆ alkoxy, or halo, and Y₆ is hydrogen.

40

5. A compound of Claim 4, in which Y₅ is halo, methoxy, or methyl.

6. A compound of Claim 4, in which at least two of X₂, X₃, X₄ and X₅ are hydrogen.

45

7. A compound of Claim 6, in which X₃ is C₁-C₆ alkyl.

8. A compound of Claim 6, in which X₃ is C₁-C₆ alkoxy.

9. A compound of Claim 1, in which m is 0 and n is 2.

50

10. A compound of claim 1, in which Y₅ is hydrogen, C₁-C₆ alkyl, C₁-C₆ alkoxy, or halo, and Y₆ is hydrogen.

11. A compound of Claim 1, in which at least two of X₂, X₃, X₄ and X₅ are hydrogen.

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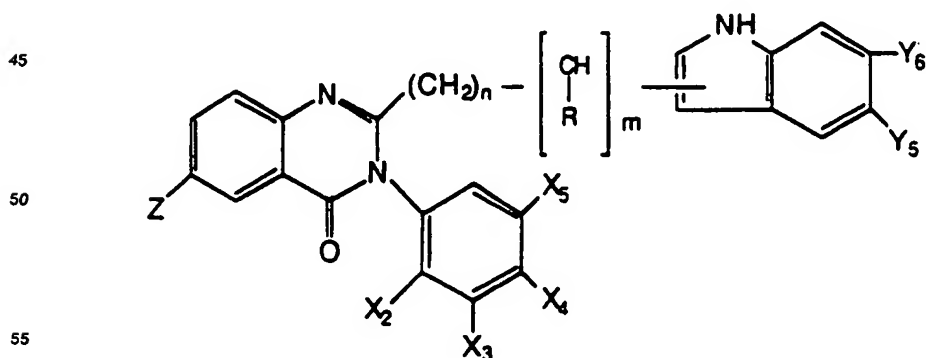
12. A compound of Claim 1, in which X₂, X₄, and X₅ are hydrogen and X₃ is C₁-C₆ alkyl.

13. A compound of Claim 1, in which X₂, X₄, and X₅ are hydrogen and X₃ is C₁-C₆ alkoxy.

14. A compound of Claim 6, in which X_2 , X_4 , and X_5 are hydrogen.
15. A compound of Claim 14, in which X_3 is isopropoxy.
- 5 16. A compound of Claim 15, in which Y_5 is hydrogen.
17. A compound of Claim 15, in which Y_5 is chloro.
18. A compound of Claim 15, in which Y_5 is fluoro.
- 10 19. A compound of Claim 15, in which Y_5 is bromo.
20. A compound of Claim 1, in which Z is halo.
- 15 21. A compound of Claim 20, in which Z is chloro.
22. A compound of Claim 21, in which m is 0 and n is 2.
23. A compound of Claim 22, in which Y_5 is chloro or bromo.
- 20 24. A compound of Claim 23, in which X_3 is C_1 - C_4 alkoxy, X_2 , X_4 , and X_5 are hydrogen.
25. A compound of Claim 1, in which m is 1 and R is selected from the group consisting of methyl, benzyl, and phenyl.
- 25 26. A compound of Claim 25, in which n is 1.
27. A compound of Claim 26, in which Y_5 is hydrogen or bromo and Y_6 is hydrogen.
- 30 28. A compound of Claim 27, in which X_3 is C_1 - C_6 alkyl or C_1 - C_6 alkoxy, and X_2 , X_4 , and X_5 are hydrogen.
29. A pharmaceutical formulation comprising as an active ingredient a compound as claimed in any one of Claims 1 to 28 associated with one or more pharmaceutically acceptable carriers, excipients, or diluents therefor.
- 35 30. A compound as claimed in any of Claims 1 to 28, for use in effecting antagonist action at the CCK and gastrin receptors.

Claims for the following Contacting State : ES

- 40 1. A process for preparing a compound of the formula



in which

n is 1 or 2 and m is 0 or 1;

R is hydrogen, C₁-C₄ alkyl, benzyl, or phenyl;

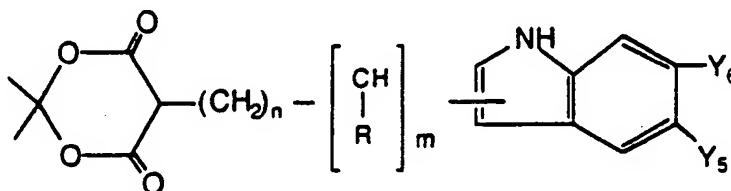
Z is hydrogen or halo;

X₂, X₃, X₄, and X₅ are independently selected from the group consisting of hydrogen, halo, trifluoromethyl, C₁-C₆ alkoxy, C₁-C₆ alkyl, C₁-C₆ alkylthio, and -NR₂R₃, in which R₂ and R₃ are independently hydrogen, C₁-C₄ alkyl, benzyl, or phenyl, or R₂ and R₃ taken together with the nitrogen atom to which they are bonded form a 5- or 6-membered ring; or

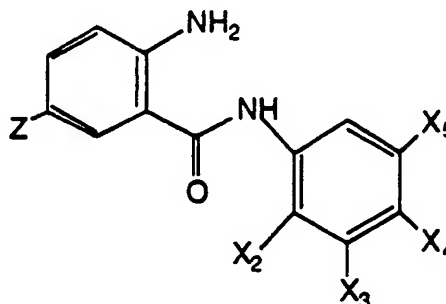
X_r and X_{r+1}, in which r is 2, 3, or 4, taken together form a divalent C₃-C₅ alkylene group or methylenedioxy; and

Y₅ and Y₆ are independently selected from the group consisting of hydrogen, C₁-C₆ alkyl, C₁-C₆ alkoxy, halo, and trifluoromethyl; which comprises

a) reacting a compound of the formula

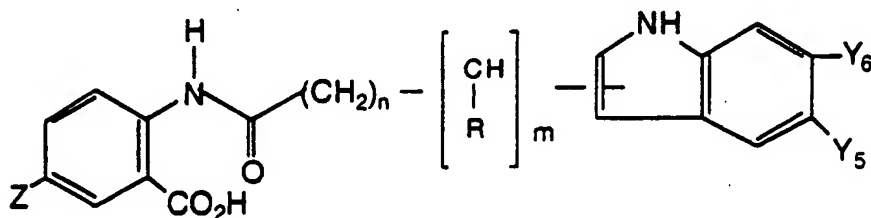


with a compound of the formula

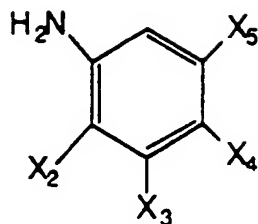


in the presence of a heteroaromatic solvent under acidic conditions; or

b) reacting a compound of the formula



with an aniline of the formula



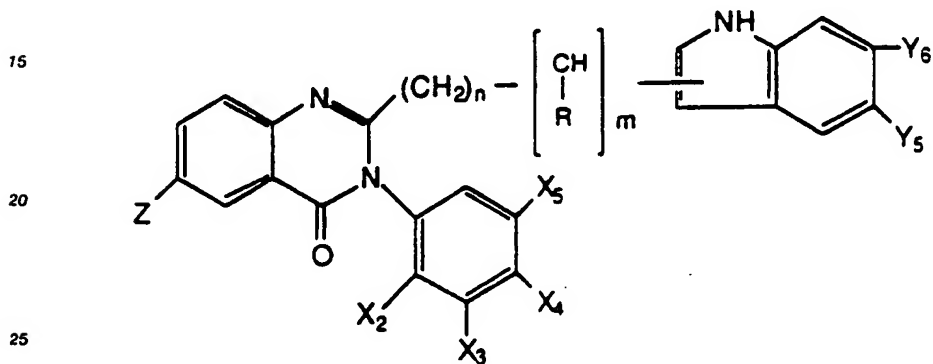
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under acidic conditions.

2. A process of Claim 1, in which Z is hydrogen.
- 15 3. A process of Claim 2, in which m is 0 and n is 2.
4. A process of Claim 3, in which Y_5 is hydrogen, $\text{C}_1\text{-C}_6$ alkyl, $\text{C}_1\text{-C}_6$ alkoxy, or halo, and Y_6 is hydrogen.
5. A process of Claim 4, in which Y_5 is halo, methoxy, or methyl.
- 20 6. A process of Claim 4, in which at least two of X_2 , X_3 , X_4 and X_5 are hydrogen.
7. A process of Claim 6, in which X_3 is $\text{C}_1\text{-C}_6$ alkyl.
- 25 8. A process of Claim 6, in which X_3 is $\text{C}_1\text{-C}_6$ alkoxy.
9. A process of Claim 1, in which m is 0 and n is 2.
10. A process of claim 1, in which Y_5 is hydrogen, $\text{C}_1\text{-C}_6$ alkyl, $\text{C}_1\text{-C}_6$ alkoxy, or halo, and Y_6 is hydrogen.
- 30 11. A process of Claim 1, in which at least two of X_2 , X_3 , X_4 and X_5 are hydrogen.
12. A process of Claim 1, in which X_2 , X_4 , and X_5 are hydrogen and X_3 is $\text{C}_1\text{-C}_6$ alkyl.
- 35 13. A process of Claim 1, in which X_2 , X_4 , and X_5 are hydrogen and X_3 is $\text{C}_1\text{-C}_6$ alkoxy.
14. A process of Claim 6, in which X_2 , X_4 , and X_5 are hydrogen.
15. A process of Claim 14, in which X_3 is isopropoxy.
- 40 16. A process of Claim 15, in which Y_5 is hydrogen.
17. A process of Claim 15, in which Y_5 is chloro.
- 45 18. A process of Claim 15, in which Y_5 is fluoro.
19. A process of Claim 15, in which Y_5 is bromo.
20. A process of Claim 1, in which Z is halo.
- 50 21. A process of Claim 20, in which Z is chloro.
22. A process of Claim 21, in which m is 0 and n is 2.
- 55 23. A process of Claim 22, in which Y_5 is chloro or bromo.
24. A process of Claim 23, in which X_3 is $\text{C}_1\text{-C}_4$ alkoxy, X_2 , X_4 , and X_5 are hydrogen.

25. A process of Claim 1, in which m is 1 and R is selected from the group consisting of methyl, benzyl, and phenyl.
26. A process of Claim 25, in which n is 1.
27. A process of Claim 26, in which Y₅ is hydrogen or bromo and Y₆ is hydrogen.
28. A process of Claim 27, in which X₃ is C₁-C₆ alkyl or C₁-C₆ alkoxy, and X₂, X₄, and X₅ are hydrogen.
29. A process for preparing a pharmaceutical formulation which comprises admixing a compound of the formula



in which

- n is 1 or 2 and m is 0 or 1;
- R is hydrogen, C₁-C₄ alkyl, benzyl, or phenyl;
- Z is hydrogen or halo;
- X₂, X₃, X₄, and X₅ are independently selected from the group consisting of hydrogen, halo, trifluoromethyl, C₁-C₆ alkoxy, C₁-C₆ alkyl, C₁-C₆ alkylthio, and -NR₂R₃, in which R₂ and R₃ are independently hydrogen, C₁-C₄ alkyl, benzyl, or phenyl, or R₂ and R₃ taken together with the nitrogen atom to which they are bonded form a 5- or 6-membered ring; or
- X_r and X_{r+1}, in which r is 2, 3, or 4, taken together form a divalent C₃-C₅ alkylene group or methylenedioxy; and
- Y₅ and Y₆ are independently selected from the group consisting of hydrogen, C₁-C₆ alkyl, C₁-C₆ alkoxy, halo, and trifluoromethyl; with one or more pharmaceutically acceptable carriers, excipients, or diluents therefor.

Patentanspruch

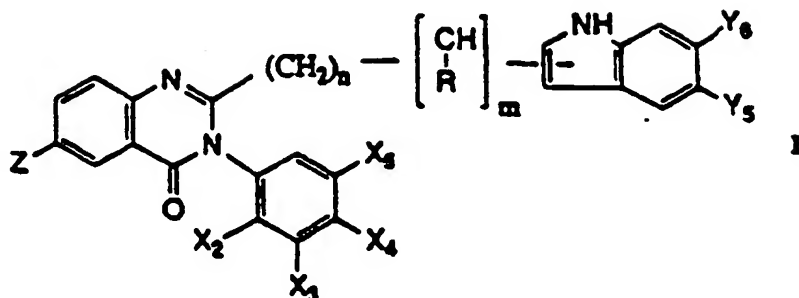
Patentanspruch für folgende Vertragsstaaten : DE, FR, GB, IT

1. Verbindung der Formel

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20 worin

 n 1 oder 2 ist und m 0 oder 1 ist; R Wasserstoff, ein C_1 - C_4 -Alkyl-, Benzyl- oder Phenylrest ist; Z Wasserstoff oder Halogen ist;

25 X_2 , X_3 , X_4 und X_5 unabhängig ausgewählt sind aus der Gruppe bestehend aus Wasserstoff, Halogen-, Trifluormethyl-, C_1 - C_6 -Alkoxy-, C_1 - C_6 -Alkyl-, C_1 - C_6 -Alkylthioestern und $-NR_2R_3$, worin R_2 und R_3 unabhängig ausgewählt sind aus Wasserstoff, C_1 - C_4 -Alkyl-, Benzyl- oder Phenylresten oder R_2 und R_3 zusammen mit dem Stickstoffatom, an das sie gebunden sind, einen fünf- oder sechsgliedrigen Ring bilden oder

30 X_r und X_{r+1} , worin r 2, 3 oder 4 ist, zusammen eine divalente C_3 - C_5 -Alkylengruppe oder einen Methylendioxyrest bilden und

Y_5 und Y_6 unabhängig ausgewählt sind aus der Gruppe bestehend aus Wasserstoff, C_1 - C_6 -Alkyl-, C_1 - C_6 -Alkoxy-, Halogen- und Trifluormethylresten; und pharmazeutisch annehmbare Salze davon.

2. Verbindung nach Anspruch 1, worin Z Wasserstoff ist.

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3. Verbindung nach Anspruch 2, worin m 0 ist und n 2 ist.4. Verbindung nach Anspruch 3, worin Y_5 Wasserstoff, ein C_1 - C_6 -Alkyl-, C_1 - C_6 -Alkoxy- oder Halogenrest ist und Y_6 Wasserstoff ist.

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5. Verbindung nach Anspruch 4, worin Y_5 Halogen, ein Methoxyrest oder ein Methylrest ist.6. Verbindung nach Anspruch 4, worin mindestens zwei der Reste X_2 , X_3 , X_4 und X_5 Wasserstoff sind.

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7. Verbindung nach Anspruch 6, worin X_3 ein C_1 - C_6 -Alkylrest ist.8. Verbindung nach Anspruch 6, worin X_3 ein C_1 - C_6 -Alkoxyrest ist.9. Verbindung nach Anspruch 1, worin m 0 und n 2 ist.

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10. Verbindung nach Anspruch 1, worin Y_5 Wasserstoff, ein C_1 - C_6 -Alkyl-, C_1 - C_6 -Alkoxy- oder Halogenrest ist und Y_6 Wasserstoff ist.11. Verbindung nach Anspruch 1, worin mindestens zwei der Reste X_2 , X_3 , X_4 und X_5 Wasserstoff sind.

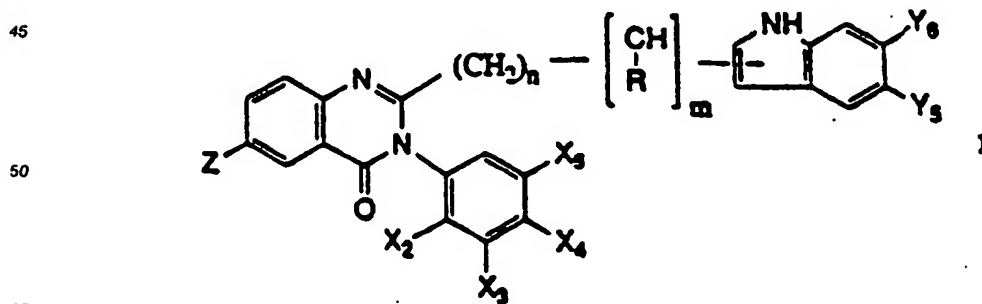
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12. Verbindung nach Anspruch 1, worin X_2 , X_4 und X_5 Wasserstoff sind und X_3 ein C_1 - C_6 -Alkylrest ist.13. Verbindung nach Anspruch 1, worin X_2 , X_4 und X_5 Wasserstoff sind und X_3 ein C_1 - C_6 -Alkoxyrest ist.

14. Verbindung nach Anspruch 6, worin X_2 , X_4 und X_5 Wasserstoff sind.
15. Verbindung nach Anspruch 14, worin X_3 ein Isopropoxyrest ist.
- 5 16. Verbindung nach Anspruch 15, worin Y_5 Wasserstoff ist.
17. Verbindung nach Anspruch 15, worin Y_5 Chlor ist.
18. Verbindung nach Anspruch 15, worin Y_5 Fluor ist.
- 10 19. Verbindung nach Anspruch 15, worin Y_5 Brom ist.
20. Verbindung nach Anspruch 1, worin Z Halogen ist.
- 15 21. Verbindung nach Anspruch 20, worin Z Chlor ist.
22. Verbindung nach Anspruch 21, worin \underline{m} 0 und \underline{n} 2 ist.
23. Verbindung nach Anspruch 22, worin Y_5 Chlor oder Brom ist.
- 20 24. Verbindung nach Anspruch 23, worin X_3 ein C_1 - C_4 -Alkoxyrest ist, X_2 , X_4 und X_5 Wasserstoff sind.
25. Verbindung nach Anspruch 1, worin \underline{m} 1 ist und R ausgewählt ist aus der Gruppe bestehend aus Methyl-, Benzyl- und Phenylresten.
- 25 26. Verbindung nach Anspruch 25, worin \underline{n} 1 ist.
27. Verbindung nach Anspruch 26, worin Y_5 Wasserstoff oder Brom ist und Y_6 Wasserstoff ist.
- 30 28. Verbindung nach Anspruch 27, worin X_3 ein C_1 - C_6 -Alkyl- oder C_1 - C_6 -Alkoxyrest ist und X_2 , X_4 und X_5 Wasserstoff sind.
29. Pharmazeutisches Präparat umfassend als aktiven Inhaltsstoff eine Verbindung nach einem der Ansprüche 1 bis 28 zusammen mit einem oder mehreren pharmazeutisch annehmbaren Trägern, Hilfsstoffen oder Verdünnungsmitteln dafür.
- 35 30. Verbindung nach einem der Ansprüche 1 bis 28 zur Verwendung, um eine antagonistische Wirkung an den CCK- und Gastrinrezeptoren zu bewirken.

40 Patentansprüche für folgenden Vertragsstaat : ES

1. Verfahren zur Herstellung einer Verbindung der Formel



worin
 \underline{n} 1 oder 2 ist und \underline{m} 0 oder 1 ist;

R Wasserstoff, ein C₁-C₄-Alkyl-, Benzyl- oder Phenylrest ist;

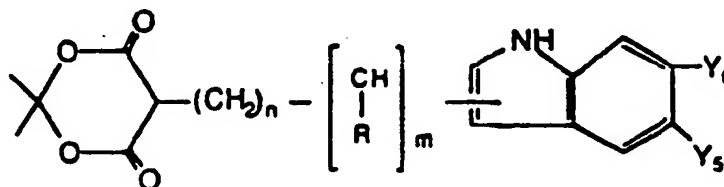
Z Wasserstoff oder Halogen n ist;

X₂, X₃, X₄ und X₅ unabhängig ausgewählt sind aus der Gruppe bestehend aus Wasserstoff, Halogen-, Trifluormethyl-, C₁-C₆-Alkoxy-, C₁-C₆-Alkyl-, C₁-C₆-Alkylthioresten und -NR₂R₃, worin R₂ und R₃ unabhängig ausgewählt sind aus Wasserstoff, C₁-C₄-Alkyl-, Benzyl- oder Phenylresten oder R₂ und R₃ zusammen mit dem Stickstoffatom, an das sie gebunden sind, einen fünf- oder sechsgliedrigen Ring bilden oder

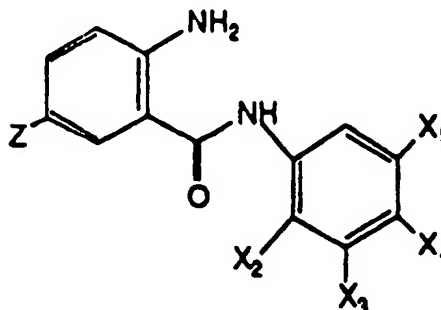
X_r und X_{r+1}, worin r 2, 3 oder 4 ist, zusammen eine divalente C₃-C₅-Alkylengruppe oder einen Methylendioxyrest bilden und

Y₅ und Y₆ unabhängig ausgewählt sind aus der Gruppe bestehend aus Wasserstoff, C₁-C₆-Alkyl-, C₁-C₆-Alkoxy-, Halogen- und Trifluormethylresten umfassend daß man

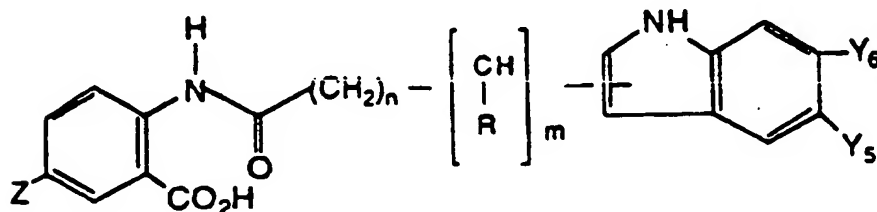
a) eine Verbindung der Formel



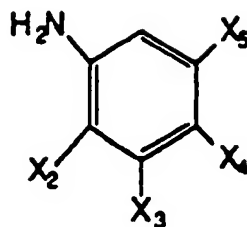
mit einer Verbindung der Formel



in Gegenwart eines heteroaromatischen Lösungsmittels unter sauren Bedingungen umgesetzt oder
b) eine Verbindung der Formel



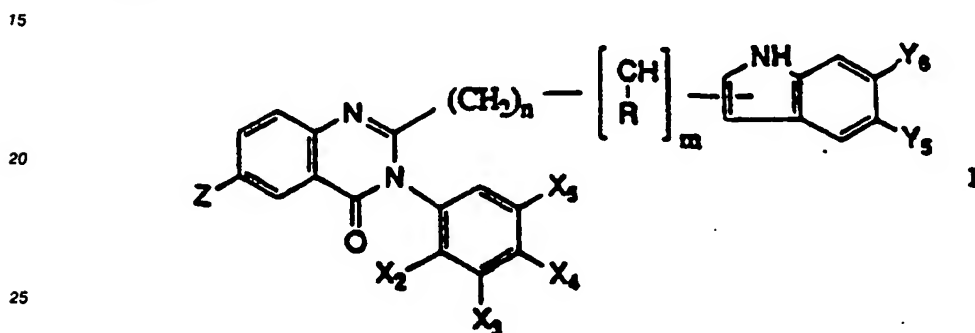
mit einem Anilin der Formel



unter sauren Bedingungen umsetzt.

2. Verfahren nach Anspruch 1, worin Z Wasserstoff ist.
- 15 3. Verfahren nach Anspruch 2, worin \underline{m} 0 und \underline{n} 2 ist.
4. Verfahren nach Anspruch 3, worin Y_5 Wasserstoff, ein C_1 - C_6 -Alkyl-, C_1 - C_6 -Alkoxy- oder Halogenrest ist und Y_6 Wasserstoff ist.
- 20 5. Verfahren nach Anspruch 4, worin Y_5 Halogen, ein Methoxyrest oder ein Methylrest ist.
6. Verfahren nach Anspruch 4, worin mindestens zwei der Reste X_2 , X_3 , X_4 und X_5 Wasserstoff sind.
- 25 7. Verfahren nach Anspruch 6, worin X_3 ein C_1 - C_6 -Alkylrest ist.
8. Verfahren nach Anspruch 6, worin X_3 ein C_1 - C_6 -Alkoxyrest ist.
9. Verfahren nach Anspruch 1, worin \underline{m} 0 und \underline{n} 2 ist.
- 30 10. Verfahren nach Anspruch 1, worin Y_5 Wasserstoff, ein C_1 - C_6 -Alkyl-, C_1 - C_6 -Alkoxy- oder Halogenrest ist und Y_6 Wasserstoff ist.
11. Verfahren nach Anspruch 1, worin mindestens zwei der Reste X_2 , X_3 , X_4 und X_5 Wasserstoff sind.
- 35 12. Verfahren nach Anspruch 1, worin X_2 , X_4 und X_5 Wasserstoff sind und X_3 ein C_1 - C_6 -Alkylrest ist.
13. Verfahren nach Anspruch 1, worin X_2 , X_4 und X_5 Wasserstoff sind und X_3 ein C_1 - C_6 -Alkoxyrest ist.
- 40 14. Verfahren nach Anspruch 6, worin X_2 , X_4 und X_5 Wasserstoff sind.
15. Verfahren nach Anspruch 14, worin X_3 ein Isopropoxyrest ist.
16. Verfahren nach Anspruch 15, worin Y_5 Wasserstoff ist.
- 45 17. Verfahren nach Anspruch 15, worin Y_5 Chlor ist.
18. Verfahren nach Anspruch 15, worin Y_5 Fluor ist.
- 50 19. Verfahren nach Anspruch 15, worin Y_5 Brom ist.
20. Verfahren nach Anspruch 1, worin Z Halogen ist.
21. Verfahren nach Anspruch 20, worin Z Chlor ist.
- 55 22. Verfahren nach Anspruch 21, worin \underline{m} 0 und \underline{n} 2 ist.
23. Verfahren nach Anspruch 22, worin Y_5 Chlor oder Brom ist.

24. Verfahren nach Anspruch 23, worin X_3 ein C_1 - C_4 -Alkoxyrest ist, X_2 , X_4 und X_5 Wasserstoff sind.
25. Verfahren nach Anspruch 1, worin m 1 ist und R ausgewählt ist aus der Gruppe bestehend aus Methyl-, Benzyl- und Phenylrest n.
26. Verfahren nach Anspruch 25, worin n 1 ist.
27. Verfahren nach Anspruch 26, worin Y_5 Wasserstoff oder Brom ist und Y_6 Wasserstoff ist.
28. Verfahren nach Anspruch 27, worin X_3 ein C_1 - C_6 -Alkyl- oder C_1 - C_6 -Alkoxyrest ist und X_2 , X_4 und X_5 Wasserstoff sind.
29. Verfahren zur Herstellung eines pharmazeutischen Präparats umfassend, daß man eine Verbindung der Formel



- worin
- n 1 oder 2 ist und m 0 oder 1 ist;
- R Wasserstoff, ein C_1 - C_4 -Alkyl-, Benzyl- oder Phenylrest ist;
- Z Wasserstoff oder Halogen ist;
- X_2 , X_3 , X_4 und X_5 unabhängig ausgewählt sind aus der Gruppe bestehend aus Wasserstoff, Halogen-, Trifluormethyl-, C_1 - C_6 -Alkoxy-, C_1 - C_6 -Alkyl-, C_1 - C_6 -Alkylthioestern und $-NR_2R_3$, worin R_2 und R_3 unabhängig ausgewählt sind aus Wasserstoff, C_1 - C_4 -Alkyl-, Benzyl- oder Phenylresten oder R_2 und R_3 zusammen mit dem Stickstoffatom, an das sie gebunden sind, einen fünf- oder sechsgliedrigen Ring bilden oder
- X_r und X_{r+1} , worin r 2, 3 oder 4 ist, zusammen eine divalente C_3 - C_5 -Alkylengruppe oder einen Methylendioxyrest bilden und
- Y_5 und Y_6 unabhängig ausgewählt sind aus der Gruppe bestehend aus Wasserstoff, C_1 - C_6 -Alkyl-, C_1 - C_6 -Alkoxy-, Halogen- und Trifluormethylresten; mit einem oder mehreren pharmazeutisch annehmbaren Trägern, Hilfsstoffen oder Verdünnungsmitteln vermischt.

Revendications

R v ndications pour l s Etats contractants suivants : DE, FR, GB, IT

1. Composé de formul

5

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20

dans laquelle

n

R

25

Z

 X_2, X_3, X_4 et X_5

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 X_r et X_{r+1} ,

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 Y_5 et Y_6

ainsi que ses sels pharmaceutiquement acceptables.

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2. Composé selon la revendication 1, dans lequel Z représente un atome d'hydrogène.

3. Composé selon la revendication 2, dans lequel m est égal à 0 et n est égal à 2.

45

4. Composé selon la revendication 3, dans lequel Y_5 représente un atome d'hydrogène, un groupe alkyle en C_1-C_6 , un groupe alcoxy en C_1-C_6 ou encore un atome d'halogène et Y_6 représente un atome d'hydrogène.

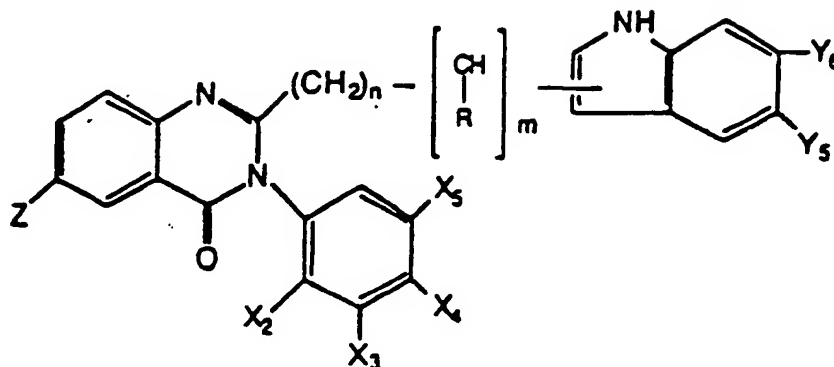
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5. Composé selon la revendication 4, dans lequel Y_5 représente un atome d'halogène, un groupe méthoxy ou un groupe méthyle.6. Composé selon la revendication 4, dans lequel au moins deux des radicaux X_2, X_3, X_4 et X_5 représentent un atome d'hydrogène.

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7. Composé selon la revendication 6, dans lequel X_3 représente un groupe alkyle en C_1-C_6 .8. Composé selon la revendication 6, dans lequel X_3 représente un groupe alcoxy en C_1-C_6 .

9. Composé selon la r vendication 1, dans lequel m est égal à 0 et n est égal à 2.



10. Composé selon la revendication 1, dans lequel Y_5 représente un atome d'hydrogène, un groupe alkyle en C_1-C_6 , un groupe alcoxy en C_1-C_6 ou encore un atome d'halogène et Y_6 représente un atome d'hydrogène.
- 5 11. Composé selon la revendication 1, dans lequel au moins deux des radicaux X_2 , X_3 , X_4 et X_5 représentent un atome d'hydrogène.
12. Composé selon la revendication 1, dans lequel X_2 , X_4 et X_5 représentent un atome d'hydrogène et X_3 représente un groupe alkyle en C_1-C_6 .
- 10 13. Composé selon la revendication 1, dans lequel X_2 , X_4 et X_5 représentent un atome d'hydrogène et X_3 représente un groupe alcoxy en C_1-C_6 .
14. Composé selon la revendication 6, dans lequel X_2 , X_4 et X_5 représentent un atome d'hydrogène.
- 15 15. Composé selon la revendication 14, dans lequel X_3 représente un groupe isopropoxy.
16. Composé selon la revendication 15, dans lequel Y_5 représente un atome d'hydrogène.
- 20 17. Composé selon la revendication 15, dans lequel Y_5 représente un atome de chlore.
18. Composé selon la revendication 15, dans lequel Y_5 représente un atome de fluor.
19. Composé selon la revendication 15, dans lequel Y_5 représente un atome de brome.
- 25 20. Composé selon la revendication 1, dans lequel Z représente un atome d'halogène.
21. Composé selon la revendication 20, dans lequel Z représente un atome de chlore.
- 30 22. Composé selon la revendication 21, dans lequel m est égal à 0 et n est égal à 2.
23. Composé selon la revendication 22, dans lequel Y_5 représente un atome de chlore ou un atome de brome.
- 35 24. Composé selon la revendication 23, dans lequel X_3 représente un groupe alcoxy en C_1-C_4 , X_2 , X_4 et X_5 représentent un atome d'hydrogène.
25. Composé selon la revendication 1, dans lequel m est égal à 1 et R est choisi parmi le groupe constitué par un groupe méthyle, un groupe benzyle et un groupe phényle.
- 40 26. Composé selon la revendication 25, dans lequel n est égal à 1.
27. Composé selon la revendication 26, dans lequel Y_5 représente un atome d'hydrogène ou un atome de brome et Y_6 représente un atome d'hydrogène.
- 45 28. Composé selon la revendication 27, dans lequel X_3 représente un groupe alkyle en C_1-C_6 ou un groupe alcoxy en C_1-C_6 et X_2 , X_4 et X_5 représentent un atome d'hydrogène.
29. Formulation pharmaceutique comprenant, comme ingrédient actif, un composé tel que revendiqué dans l'une quelconque des revendications 1 à 28, en association avec un ou plusieurs supports, excipients
- 50 ou diluants pharmaceutiquement acceptables pour le composé.
30. Composé tel que revendiqué dans l'une quelconque des revendications 1 à 28, à utiliser pour la mise en œuvre d'une action antagoniste aux récepteurs de la CCK et de la gastrine.

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R revendications pour l'Etat contractant suivant : ES

1. Procédé pour préparer un composé de formule

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15

20

dans laquelle

n

R

25

Z

X₂, X₃, X₄ et X₅

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X_r et X_{r+1},

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Y₅ et Y₆

a) faire réagir un composé de formule

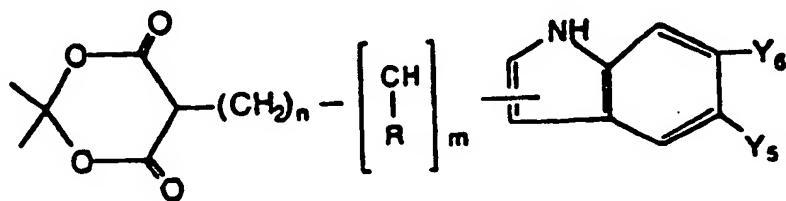
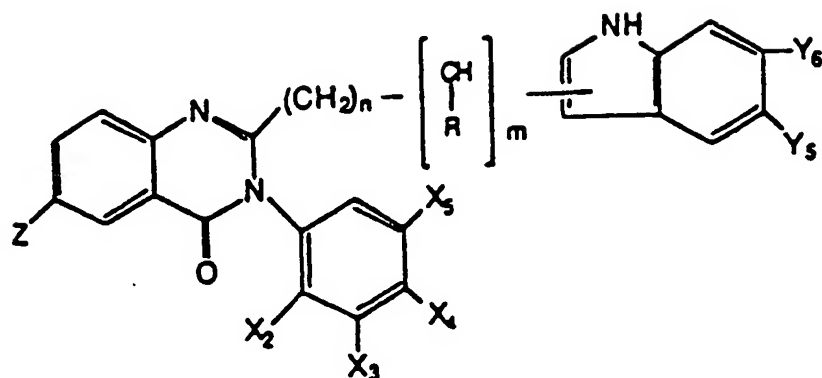
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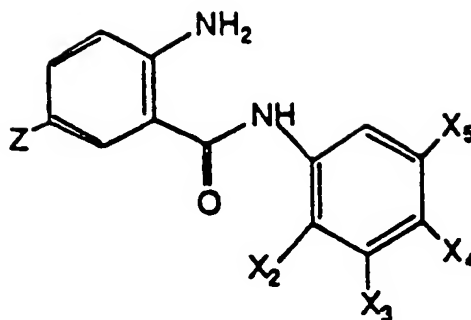
avec un composé de formule

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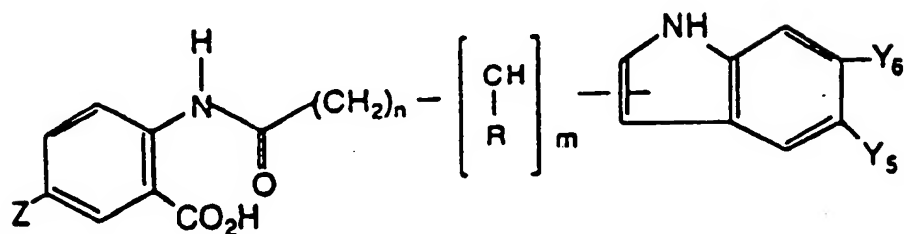


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en présence d'un solvant hétéro-aromatique dans des conditions acides; ou
b) faire réagir un composé de formule

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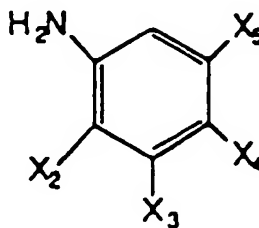
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avec une aniline de formule

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dans des conditions acides.

2. Procédé selon la revendication 1, dans lequel Z représente un atome d'hydrogène.

3. Procédé selon la revendication 2, dans lequel m est égal à 0 et n est égal à 2.

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4. Procédé selon la revendication 3, dans lequel Y5 représente un atome d'hydrogène, un groupe alkyle en C₁-C₆, un groupe alcoxy en C₁-C₆ ou encore un atome d'halogène et Y6 représente un atome d'hydrogène.

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5. Procédé selon la revendication 4, dans lequel Y5 représente un atome d'halogène, un groupe méthoxy ou un groupe méthyle.

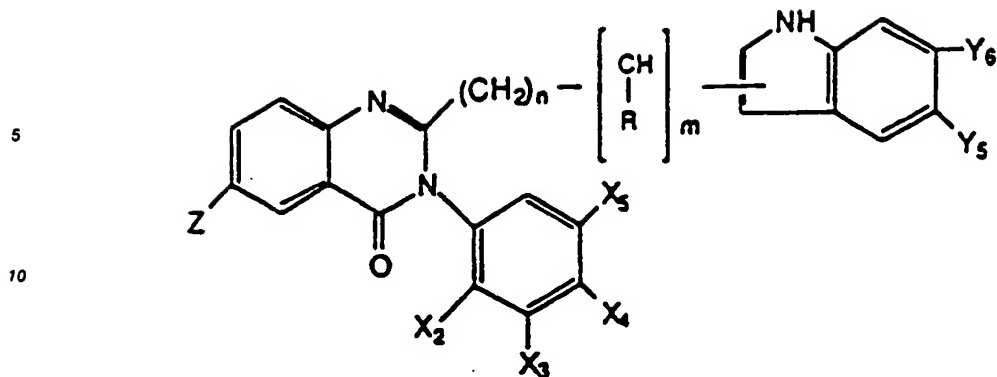
6. Procédé selon la revendication 4, dans lequel au moins deux des radicaux X₂, X₃, X₄ et X₅ représentent un atome d'hydrogène.

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7. Procédé selon la revendication 6, dans lequel X₃ représente un groupe alkyle en C₁-C₆.

8. Procédé selon la revendication 6, dans lequel X₃ représente un groupe alcoxy en C₁-C₆.

9. Procédé selon la revendication 1, dans lequel m est égal à 0 et n est égal à 2.
10. Procédé selon la revendication 1, dans lequel Y₅ représente un atome d'hydrogène, un groupe alkyle en C₁-C₆, un groupe alcoxy en C₁-C₆ ou encore un atome d'halogène et Y₆ représente un atome d'hydrogène.
11. Procédé selon la revendication 1, dans lequel au moins deux des radicaux X₂, X₃, X₄ et X₅ représentent un atome d'hydrogène.
12. Procédé selon la revendication 1, dans lequel X₂, X₄ et X₅ représentent un atome d'hydrogène et X₃ représente un groupe alkyle en C₁-C₆.
13. Procédé selon la revendication 1, dans lequel X₂, X₄ et X₅ représentent un atome d'hydrogène et X₃ représente un groupe alcoxy en C₁-C₆.
14. Procédé selon la revendication 6, dans lequel X₂, X₄ et X₅ représentent un atome d'hydrogène.
15. Procédé selon la revendication 14, dans lequel X₃ représente un groupe isopropoxy.
16. Procédé selon la revendication 15, dans lequel Y₅ représente un atome d'hydrogène.
17. Procédé selon la revendication 15, dans lequel Y₅ représente un atome de chlore.
18. Procédé selon la revendication 15, dans lequel Y₅ représente un atome de fluor.
19. Procédé selon la revendication 15, dans lequel Y₅ représente un atome de brome.
20. Procédé selon la revendication 1, dans lequel Z représente un atome d'halogène.
21. Procédé selon la revendication 20, dans lequel Z représente un atome de chlore.
22. Procédé selon la revendication 21, dans lequel m est égal à 0 et n est égal à 2.
23. Procédé selon la revendication 22, dans lequel Y₅ représente un atome de chlore ou un atome de brome.
24. Procédé selon la revendication 23, dans lequel X₃ représente un groupe alcoxy en C₁-C₄, X₂, X₄ et X₅ représentent un atome d'hydrogène.
25. Procédé selon la revendication 1, dans lequel m est égal à 1 et R est choisi parmi le groupe constitué par un groupe méthyle, un groupe benzyle et un groupe phényle.
26. Procédé selon la revendication 25, dans lequel n est égal à 1.
27. Procédé selon la revendication 26, dans lequel Y₅ représente un atome d'hydrogène ou un atome de brome et Y₆ représente un atome d'hydrogène.
28. Procédé selon la revendication 27, dans lequel X₃ représente un groupe alkyle en C₁-C₆ ou un groupe alcoxy en C₁-C₆ et X₂, X₄ et X₅ représentent un atome d'hydrogène.
29. Procédé pour préparer une formulation pharmaceutique, qui comprend le fait de mélanger un composé de formule



dans laquelle

n

est égal à 1 ou 2 et m est égal à 0 ou 1;

R

représente un atome d'hydrogène, un groupe alkyle en C₁-C₄, un groupe benzyle ou un groupe phényle;

Z

représente un atome d'hydrogène ou un atome d'halogène;

X₂, X₃, X₄ et X₅

sont choisis, indépendamment l'un de l'autre, parmi le groupe constitué par un atome d'hydrogène, un atome d'halogène, un groupe trifluorométhyle, un groupe alcoxy en C₁-C₆, un groupe alkyle en C₁-C₆, un groupe alkyl(en C₁-C₆)thio et un groupe -NR₂R₃ dans lequel R₂ et R₃ représentent, indépendamment l'un de l'autre, un atome d'hydrogène, un groupe alkyle en C₁-C₄, un groupe benzyle ou un groupe phényle, ou bien R₂ et R₃, pris ensemble avec l'atome d'azote auquel ils sont liés, forment un noyau penta- ou hexagonal; ou

X_r et X_{r+1},

où r représente 2, 3 ou 4, pris ensemble, forment un groupe alkylène en C₃-C₅ divalent ou un groupe méthylènedioxy; et

Y₅ et Y₆

sont choisis, indépendamment l'un de l'autre, parmi le groupe constitué par un atome d'hydrogène, un groupe alkyle en C₁-C₆, un groupe alcoxy en C₁-C₆, un atome d'halogène et un groupe trifluorométhyle, avec un ou plusieurs supports, excipients ou diluants pharmaceutiquement acceptables pour le composé.